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Dose dependence of magnetism in Co-doped TiO₂

N. Akdogan^{a,*}, A. Nefedov^a, A. Westphalen^a, H. Zabel^a,
R.I. Khaibullin^b, L.R. Tagirov^{b,c}

^a *Institut für Experimentalphysik/Festkörperphysik, Ruhr-Universität Bochum, D-44780 Bochum, Germany*

^b *Kazan Physical-Technical Institute of RAS, 420029 Kazan, Russia*

^c *Kazan State University, 420008 Kazan, Russia*

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Abstract

The dose dependence of magnetism in Co-implanted TiO₂ rutile was investigated using the magneto-optic Kerr effect method at room temperature. The (100)- and (001)-oriented single-crystalline rutile TiO₂ plates were used as substrates and implanted by Co ions with a varying dose range of $0.25\text{--}1.50 \times 10^{17}$ ions/cm². We observed paramagnetic behaviour for the low dose doped samples, but obtain clear hysteretic-like behaviour for intermediate and high dose doped samples. For the intermediate implantation doses of Co, ferromagnetic behaviour can be explained by the F-center (oxygen vacancies) exchange mechanism, while for the highest implantation dose, in addition to Co substitution on Ti sites, formation of Co nanoclusters may take place within the TiO₂ irradiated region.

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The discovery of room temperature ferromagnetism in Co-doped anatase TiO₂ [1] has generated much interest in the Co:TiO₂ system as a potential oxide-based diluted magnetic semiconductor (DMS) [2–7]. In spite of rather extensive studies in this field, the origin of the observed ferromagnetism is still unclear. Several investigations explained that cobalt ions in thin TiO₂ films exist in a +2 oxidation state, consistent with ferromagnetism that originates from Co substitution on Ti sites [2], while in other publications the origin of ferromagnetism is reported to be due to the precipitation of cobalt metal clusters [8,9]. Recently, we have reported room temperature ferromagnetism and in-plane magnetic anisotropy of single-crystalline rutile

* Corresponding author. Tel.: +49 0 234 32 23626.

E-mail address: numan.akdogan@rub.de (N. Akdogan).